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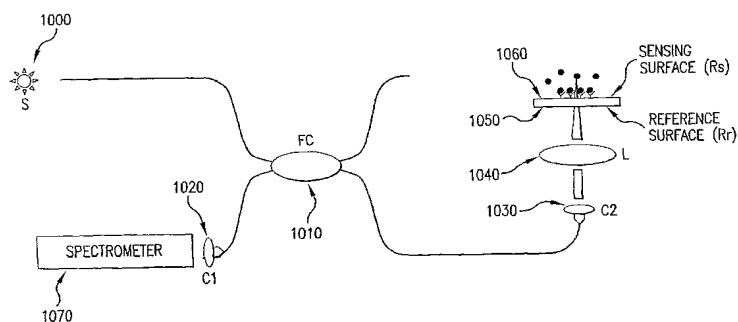
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(54) Title: ARRANGEMENTS, SYSTEMS AND METHODS CAPABLE OF PROVIDING SPECTRAL-DOMAIN OPTICAL COHERENCE REFLECTOMETRY FOR A SENSITIVE DETECTION OF CHEMICAL AND BIOLOGICAL SAMPLE



(57) Abstract: Systems, arrangements and methods for a molecular recognition are provided. For example, a particular radiation having wavelength that varies over time and/or a spectral width that is greater than 10nm can be provided. For example, at least one first electro-magnetic radiation can be provided to at least one sample, and at least one second electro-magnetic radiation may be provided to a reference, with both the first and second electro-magnetic radiations being part of the particular radiation. Further, the interference between a third electro-magnetic radiation (associated with the first electro-magnetic radiation) and a fourth electro-magnetic radiation (associated with the second electro-magnetic radiation) can be detected. A change in a thickness of at least one portion of the sample based on the interference can be determined.



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**ARRANGEMENTS, SYSTEMS AND METHODS CAPABLE OF PROVIDING
SPECTRAL-DOMAIN OPTICAL COHERENCE REFLECTOMETRY FOR A
SENSITIVE DETECTION OF CHEMICAL AND BIOLOGICAL SAMPLE**

5 CROSS-REFERENCE TO RELATED APPLICATION(S)

This application is based upon and claims the benefit of priority from U.S. Patent Application Serial No. 60/680,947, filed May 13, 2005, the entire disclosure of which is incorporated herein by reference.

10 STATEMENT REGARDING FEDERALLY SPONSORED RESEARCH

The invention was made with the U.S. Government support under Contract No. RO1 EY014975 and RO1RR019768 awarded by the National Institute of Health, and Contract No. F49620-021-1-0014 awarded by the Department of Defense. Thus, the U.S. Government has certain rights in the invention.

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FIELD OF THE INVENTION

The present invention relates to methods and apparatus for a molecular recognition. More particularly, the present invention relates to detection arrangements, systems and methods for a molecular binding on a sensing surface and the presence of molecules in channels.

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BACKGROUND OF THE INVENTION

Real-time detection of minute traces of molecules (e.g., pesticides, viruses, and organic toxins) is important in various applications such as medical diagnostics, environmental monitoring, and homeland security. For example, there is a need for providing a highly sensitive detection methods of viruses, as well as processes that provide an early detection of chemicals and pathogens (e.g., explosives, anthrax) which could trigger a corrective action. Such methods may be important in a broad range of, e.g., medical and environmental applications and bio-defense.

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Such exemplary detection has been conducted by fluorescent (as described in D. W. Pierce et al., "Imaging individual green fluorescent proteins," Nature, 1997, Vol. 388, pp. 338 et seq.) and using certain radioactive methods. Even

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though these label-based techniques could potentially achieve single molecular level detection, an additional specimen preparation is needed to be performed therefor, which is costly in time and may affect the molecules of interest.

Label-free detection techniques such as surface plasmon resonance (SPR) sensors (as described in J. Homola et al., "Surface plasmon resonance sensors: review," *Sensors and Actuators B*, 1999, Vol. 54, pp. 3-15) and quartz crystal microbalances (QCM) arrangements (as described in G. Kleefisch et al., "Quartz microbalance sensor for the detection of Acrylamide," *Sensors*, 2004, Vol. 4, pp. 136-146) provide an indication of a physical absorption of molecules on a sensor surface.

The SPR sensor generally exploits the change of the SPR angle due to the alteration of refractive index at a metal-dielectric interface upon the protein absorption. However, this sensor may review a large amount of molecules, since its lateral resolution may not be reduced without loss of sensitivity (as described in C. Berger et al., "Resolution in surface plasmon microscopy," *REVIEW OF SCIENTIFIC INSTRUMENTS*, 1994, Vol. 65, pp. 2829-2836). QCM techniques also utilize the shift of resonance frequency due to the effective mass increase upon the protein binding. In addition to the needed large amount of molecules, the QCM detection method needs to operate in a dry environment, preferably in a vacuum, because the damping in aqueous environment likely deteriorates the sensitivity.

Several methods based on micro-fabrication techniques have been (as provided in P. Burg et al., "Suspended microchannel resonators for biomolecular detection," *Applied Physics Letters*, 2003, Vol. 83(13), pp. 2698-2700; and W. U. Wang et al., "Label-free detection of small-molecule-protein interactions by using nanowire nanosensors," *PROCEEDINGS OF THE NATIONAL ACADEMY OF SCIENCES OF THE UNITED STATES OF AMERICA*, 2005. 102: p. 3208-3212), attempted to address the above-described deficiencies. Such methods could potentially achieve sensitive detection for label-free species, but the fabrication techniques (e.g., e-beam lithography, electron beam evaporation, and chemical vapor deposition) are complicated and expensive, and the sensing units that use such techniques are likely directly coupled to micro-fluidic devices, limiting their utility for various diagnostic applications.

A spectral domain optical coherence reflectometry (SD-OCR) technique is an optical ranging procedure which is capable of measuring depth-resolved phase information with a sub-nanometer thickness sensitivity. For example, a thickness change can be an optical thickness change, a refractive index change, and/or a physical thickness change. Detailed descriptions on SD-OCR and demonstration of sub-nanometer sensitivity are provided in International Patent Application PCT/US03/02349 and described in C. Joo et al., "Spectral-domain optical coherence phase microscopy for quantitative phase-contrast imaging," Optics Letters, 2005, Vol. 30, pp. 2131-2133; and B.C. Nassif et al., "In vivo human retinal imaging by ultrahigh-speed spectral domain optical coherence tomography," Optics Letters, 2004, Vol. 29, pp. 480-482.

OBJECTS AND SUMMARY OF THE INVENTION

One of the objects of the present invention is to overcome certain deficiencies and shortcomings of the prior art systems (including those described herein above), and implement exemplary SD-OCR techniques as shall be described in further detail below. This can be done by implementing arrangements, systems and methods which utilize SD-OCR techniques (e.g., SD-OCR arrangements, systems and methods). Another object of the present invention is to utilize systems, arrangements and methods and apply SD-OCR techniques to obtain a highly sensitive detection of label-free chemical and biological species (e.g., anatomical samples).

For example, exemplary embodiments of the system, arrangement and method according to the present invention can be provided for label-free chemical and biological species. The exemplary embodiments can utilize a coherence gating of low-coherence interferometry to identify the interference signal of interest, and measures the phase alteration of that signal for molecular absorption/removal at a surface or concentration measurement in the channels. For molecular binding on a sensing surface, these exemplary embodiments can permit an examination of molecular interactions on a micron-sized area, and thus can be extended to monitoring a large number of activated sites in parallel on a two-dimensional surface in disposable arrays, and can be adapted for the detection of new chemical and biological species by including an active binding site into the micro arrays.

Therefore, systems, arrangements and methods for a molecular (e.g., for a molecular binding on a sensing surface and the presence of molecules in channels) are provided. For example, a particular radiation having wavelength that varies over time and/or a spectral width that is greater than 10nm can be provided.

5 For example, at least one first electro-magnetic radiation can be provided to at least one sample, and at least one second electro-magnetic radiation may be provided to a reference, with both the first and second electro-magnetic radiations being part of the particular radiation. Further, the interference between a third electro-magnetic radiation (associated with the first electro-magnetic radiation) and a fourth electro-

10 magnetic radiation (associated with the second electro-magnetic radiation) can be detected. A change in a thickness of at least one portion of the sample based on the interference can be determined.

According to another exemplary embodiment of the present invention, the first and second radiations can share a common path. The sample can include a

15 plurality of samples, and the change in the thickness of the at least one portion of each of the samples may be determined simultaneously. The change in the thickness of the at least one portion of the at least one samples may be determined simultaneously at different locations along and/or perpendicular to a beam path of the first electro-magnetic radiation. The change in the thickness may also be determined

20 simultaneously along different locations along a beam path of the first electro-magnetic radiation. The first electro-magnetic radiation may be scanned over a surface of the sample at a plurality of locations thereon.

According to still another exemplary embodiment of the present invention, the portion of the sample may be coated with particular molecules that are

25 designed to associate with or dissociate from to further molecules. The change of the thickness may be associated with an association or a dissociation of the particular molecules. The particular molecules may have an affinity to bind to the further molecules that are different from the particular molecules. The portion may include a plurality of portions. For example, a first set of the particular molecules may have an

30 affinity to bind to a first portion of the portions, and a second set of the particular molecules can have an affinity to bind to a second portion of the of portions. The first and second sets may be different from one another.

In a further exemplary embodiment of the present invention, the sample can have multiple layers therein and/or may be disposable. The sample can be a micro-fluidic arrangement. The change of the thickness of the portion of the sample can be an optical thickness change and/or a physical thickness change and/or a refractive index change. The thickness change can be associated with a concentration of molecules of on and/or in the portion of the sample. The thickness can change as a function of wavelength that is associated with types of molecules of on and/or in the portion of the sample. The first electro-magnetic radiation may have a cross-section of a beam on and/or in the portion of the sample has a size that can be as small as a diffraction-limited size (e.g., 10 μ m). The thickness can be determined by (i) transforming the interference into first data which is in a complex format, (ii) determining an absolute value associated with the first data to generate second data, (iii) identifying particular locations of the portion as a function of the second data, (iv) determining a phase associated with the first data to generate third data, and (v) associating the change of the thickness with the third data. Further, the interference may be Fourier transformed to generate the first data.

These and other objects, features and advantages of the present invention will become apparent upon reading the following detailed description of embodiments of the invention, when taken in conjunction with the appended claims.

BRIEF DESCRIPTION OF THE DRAWINGS

Further objects, features and advantages of the invention will become apparent from the following detailed description taken in conjunction with the accompanying figures showing illustrative embodiments of the invention, in which:

Figure 1 is a diagram of an exemplary embodiment of an SD-OCR biosensing arrangement in accordance with the present invention;

Figure 2a is diagram of an exemplary usage of the exemplary arrangement of Figure 1 for a measurement of a molecular interaction at a particular point in time in accordance with the present invention;

Figure 2b is diagram of the exemplary usage of the exemplary arrangement of Figure 1 for the measurement of the molecular interaction at a subsequent point in time in accordance with the present invention;

Figure 2c is diagram of the exemplary usage of the exemplary arrangement of Figure 1 for the measurement of the molecular interaction at a still subsequent point in time in accordance with the present invention;

Figure 3 is a diagram of the exemplary embodiment of the SD-OCR
5 arrangement which is illustrated as performing a SD-OCR depth-resolved measurement of the molecular interaction;

Figure 4 is an exemplary operational measurement in accordance with an exemplary embodiment of the present invention using the SD-OCR biosensing arrangements of Figure 1 and/or Figure 3 and/or the arrangements described in
10 International Patent Application PCT/US03/02349 to measure the depth-resolved information, e.g., at all interfaces simultaneously, and graph associated therewith which shows the outputs thereof;

Figure 5 is an operational measurement diagram in accordance with an exemplary embodiment of the present invention using the SD-OCR biosensing
15 arrangement of Figure 1 and/or Figure 3 and/or the arrangements described in International Patent Application PCT/US03/02349 which provides a multi-channel detection of the molecular interaction, and graph associated therewith which shows the outputs thereof;

Figure 6a is an operational measurement in accordance with an
20 exemplary embodiment of the present invention using the SD-OCR biosensing arrangement of Figure 1 and/or Figure 3 and/or the arrangements described in International Patent Application PCT/US03/02349 for monitoring a phase in the interference between reflected beams from top and bottom surfaces of a microfluidic device as a function of time;

Figure 6b is an operational measurement in accordance with an
25 exemplary embodiment of the present invention using the SD-OCR biosensing arrangement of Figure 1 and/or Figure 3 and/or the arrangements described in International Patent Application PCT/US03/02349 to performing the concentration monitoring procedure of Figure 6a with the aid of a galvanometer beam scanner;

Figure 7 is a graph illustrating exemplary Subsequent bBSA-
30 streptavidin bindings measured by the exemplary SD-OCR biosensing arrangement according to the present invention;

Figure 8a is a graph showing results of an exemplary controlled bBSA-streptavidin binding measurement illustrating an increase in a thickness at a bBSA-functionalized sensor surface;

Figure 8b is a graph showing results of an exemplary controlled bBSA-streptavidin binding measurement which illustrates that no increase in the thickness was observed in a non-functionalized surface;

Figure 9a is a graph showing an exemplary change of a cover slip thickness at a particular HF concentration in accordance with the present invention;

Figure 9b is a graph showing an exemplary change of an etching rate at different HF concentrations in accordance with the present invention;

Figure 10 is an exemplary graph of an image of a photosynthetic protein layer generated using the arrangement and method in accordance with the present invention; and

Figure 11 is a flow diagram of an exemplary embodiment of the method according to the present invention.

Throughout the figures, the same reference numerals and characters, unless otherwise stated, are used to denote like features, elements, components or portions of the illustrated embodiments. Moreover, while the present invention will now be described in detail with reference to the figures, it is done so in connection with the illustrative embodiments.

DETAILED DESCRIPTION OF EXEMPLARY EMBODIMENTS OF INVENTION

An exemplary embodiment of a fiber-based SD-OCR system according to the present invention is depicted as a diagram in Figure 1. For example, as shown in Figure 1, the system can include a broadband light source (1000) which may be configured to illuminate an interferometer (1010) such as a 2×2 fiber coupler, and the beam may be focused onto a sensing surface with a diffraction limited spot size. The sensing surface can be a protein/DNA chip or a part of a micro-fluidic device. The reflected beams from the interfaces of the sensing surface 1060 (and a glass 1050) can be re-coupled to the interferometer to produce an interference signal at the detection arm. At the spectrometer (1070), the signal related to the interference may be expressed as:

$$I(k) = 2\sqrt{R_r R_s(z)} S(k) \cos(2k\Delta p), \quad (1)$$

where k is the wave number, z is the geometrical distance, and R_r and $R_s(z)$ represent the reference reflectivity and measurement reflectivity at depth z , respectively. $S(k)$ is the power spectral density of the source, and Δp is the optical path length

- 5 difference between the reference and measurement beams. A complex-valued depth information $F(z)$ is obtained by a discrete Fourier transform of Equation (1) with respect to $2k$, so the intensity and phase at depth z can be obtained as:

$$I(z) = |F(z)|^2, \quad (2)$$

$$\phi(z) = \tan^{-1} \left[\frac{\text{Im}(F(z))}{\text{Re}(F(z))} \right] = 2 \frac{2\pi}{\lambda_0} \Delta p(z), \quad (3)$$

- 10 where λ_0 is the center wavelength of the source. The depth-resolved intensity information in Equation (2) is used to locate a specific interference signal of interest, and the phase (or thickness) alteration at that signal is monitored in real-time for molecular recognition. Indeed, the spectrometer (1070) can measure power spectrum of the interference between the reference (bottom surface of a glass 1050) and the
15 molecule-coupled sensing surface or slide (1060). The system also can include collimators (C1: 1020, C2: 1030), focusing lens (L: 1040) and spectrometer (1070).

For example, to perform an exemplary molecular absorption detection, exemplary probe molecules at the sensing surface can be immobilized or patterned via known protocols (as described in BIACORE Getting Started. 1998, Biacore AB).

- 20 One of the ways to perform this can be by immersing the sensor surface in a high concentration solution of the probe molecules for several hours, and then rinse it with a Phosphate Buffered Saline (PBS) solution. In terms of patterning an array of probe molecules, this can be done by employing a micro-contact printing technique (as described in A. Bernard et al., "Microcontact printing of proteins," Advanced
25 Materials, 2000, Vol. 12, pp. 1067-1070), in which a polydimethylsiloxane (PDMS) stamp containing protein is brought into contact with the surface for physical absorption. After the sensor surface is activated with the probes, the analytes may be introduced to the sensing surface, as shown in Figures 2a-2c which illustrates an

exemplary measurement of the molecular interaction using the exemplary system of Figure 1. For example, probe molecules (2020) can be immobilized on the sensing surface (2010), and the molecules of interest (2030) can be introduced. As the analytes interact and bind to the probe molecules, the thickness at the sensor surface changes, and the reflection from the layer of bound molecules leads to a phase alteration in the interference signal being measured. In other words, as the molecules bind to the probe molecules, the phase change can be detected in real-time. This exemplary change is utilized to study the affinity of the analytes to the probe molecules and the kinetics associated with the interaction.

Exemplary embodiments of the system, arrangement and method according to the present invention can also provide a depth-resolved detection of molecular interactions, as shown in Figure 3 which illustrates another exemplary embodiment of the SD-OCR arrangement which can perform a SD-OCR depth-resolved measurement of the molecular interaction. As shown in Figure 3, the mirror (M: 3080) can be provided in the reference path, and the spectrometer (3090) may measure the power spectrum of the interference between the reflection from the reference mirror (M: 3080) and the reflections from the molecule-coupled glass slides (3050, 3060). In particular, this exemplary arrangement of Figure 3 may further include a broadband light source (S: 3000), a 2×2 fiber coupler (FC: 3010), collimators (C1: 3020, C2: 3030, C3: 3070), a focusing lens (L: 3040), molecule-coupled glass slides (3050, 3060), and spectrometer (3090). For example, the interference can be measured between the reflected beam from the stationary mirror and the beams from the interfaces of the multilayer device is measured.

Figure 4 illustrates an exemplary operational measurement in accordance with an exemplary embodiment of the present invention using the SD-OCR biosensing arrangement of Figure 1 and/or Figure 3 and/or the arrangements described in International Patent Application PCT/US03/02349 to measure the depth-resolved information, e.g., at most or all interfaces simultaneously, and a graph associated therewith which shows the outputs thereof. For example, the electromagnetic radiation or light can be projected via one or more lenses L (4000), and molecule-coupled sensor surfaces (4010, 4020) shown in this figure can be activated with different molecules. An exemplary depth-resolved measurement based on these

surfaces (4010, 4020) may indicate different affinities of molecules of interest with the immobilized molecules A and B. The intensity information can be used to identify each sensor surface (3050, 3060) shown in Figure 3, and the phase of each such sensor surface can be monitored in real-time for analyzing kinetics of the same or similar analytes for the difference (probe) molecules, for example as shown in Figure 4.

An exemplary embodiment of a high-throughput multi-channel detection of molecular bindings is possible via micro arrays of probe molecules as shown in the diagram and graph of Figure 5. As shown in Figure 5, which illustrates a galvanometer scanning mirror (GM: 5000), a focusing lens (L: 5010), and a multi-molecule coupled glass slide (5020), a sensor surface of the slide (5020) can be patterned with small features (1~10 μm) of different probes, after which the free surface is saturated with inert proteins. As the molecules of interest, or analytes, are introduced, the probe beam scans across the sensing surface to monitor and measure molecular interactions in each of probe (or activation) sites in real-time. Since non-specific protein-protein binding (cross reactivity) is common to the entire sensor surface, it can be cancelled out by examining the entire sensor surface and by comparing the change in probe (or activated) regions with that of non-activated regions.

In addition to the detection of molecular absorption on a sensing surface, exemplary embodiments of the system, arrangement and method according to the present invention can also be used for measuring the amount (or concentration) of the free molecules in a fluidic channel. For example, the presence of the free molecules in a solution can change the effective refractive index in the channel, which may alter the phase in the interference between the reflected beams from the top and bottom surfaces of the channel. Figures 6a and 6b show operational illustrations of two exemplary depictions of such concepts, and include at least one focusing lens (L: 6000), a microfluidic device (6010), and a galvanometer beam scanner (GM: 6030). In Figure 6a, the phase in the interference between the top and bottom walls of the fluidic channel is measured or monitored at one or more specific locations as a function of time, and the introduction of the molecules in the channel increases the phase measurement. Through an appropriate calibration, the exemplary embodiments

of the present invention can be used to quantify the concentration level of the solution. Figure 6b shows an operational diagram of how two different molecules diffuse in a fluidic channel. As provided in this drawing, the probe beam scans across the fluidic channel to measure the spatial phase distribution, as the molecules diffuse.

5 As the molecules are flowing into the channel, e.g., between surfaces of the microfluidic device (6010), the phase change can be induced, which may indicate the change in the molecule concentration. For the diffusion measurement, the probe beam scans across the channel, and measure the spatial phase distribution caused by diffusion of these molecules. This measurement can be useful to quantify diffusion

10 rate and binding affinity of label-free species for a given environment.

SUPPORTING DATA

I. Measurement of biotin and streptavidin interaction

As a preliminary demonstration of the implementation of the

15 exemplary embodiments of the present invention, the interaction between biotin and streptavidin at a sensor surface was measured as provided in Figure 7, which shows a graph 7010 of exemplary Subsequent bBSA-streptavidin bindings measured by the exemplary SD-OCR biosensing arrangement according to the present invention. The interior channel of a micro-fluidic device was activated with biotinylated bovine

20 serum albumin (bBSA), and several experiments were conducted to detect the subsequent bBSA-streptavidin bindings. Initially, introduction of PBS solution did not change the thickness at the sensing surface, but the noticeable change was observed after the streptavidin solution (1 μ M) was injected into the exemplary device, due to the binding of the streptavidin to the immobilized bBSA layer. As shown in Figure 7,

25 the thickness remained constant after all the binding sites of bBSA were occupied by the streptavidin. The subsequent introduction of PBS solution did not change the thickness measurement. However, when the bBSA solution was flowed in again (3 μ M), a further increase in the thickness was observed, which can be because the injection of streptavidin restored the ability to bind bBSA in the channel, as illustrated

30 by bBSA-streptavidin multi-layer formation. The subsequent introduction of the buffer solution did not change the signal, but when switched back to bBSA solution, a further thickness increase was observed.

Control experiments with lower concentration of streptavidin solution (250 nM) were also conducted as provided in Figures 8a and 8b. For example, Figure 8a shows a graph 8010 providing exemplary results of an exemplary controlled bBSA-streptavidin binding measurement illustrating an increase in an thickness at a bBSA-functionalized sensor surface. Figure 8b shows a graph 8020 of exemplary results of the exemplary controlled bBSA-streptavidin binding measurement which illustrates that no increase in the thickness was observed in a non-functionalized surface. As shown in these drawings, the channel of a micro-fluidic device was functionalized with bBSA, and the streptavidin was introduced into the channel. The thickness increase was observed due to the binding of the streptavidin with slower rate, compared to a previous measurement. However, in the case of non-functionalized sensing surface, the thickness did not change, as shown in Figure 8b, which demonstrates specific binding nature of streptavidin with biotin.

II. Detection of SiO₂ etching

A flow diagram of the exemplary embodiment of the method according to the present invention is shown in Figure 11. For example, a particular radiation having wavelength that varies over time and/or a spectral width that is greater than 10nm can be provided by a source arrangement (step 110). Indeed, a first electro-magnetic radiation can be provided to sample and a second electro-magnetic radiation may be provided to a reference (both being part of particular radiation) as provided in step 120. Next, the interference between a third electro-magnetic radiation (associated with the first electro-magnetic radiation) and a fourth electro-magnetic radiation (associated with the second electro-magnetic radiation) can be detected in step 130. Further, a change in a thickness of at least one portion of the sample based on the interference can be determined in step 140.

The exemplary embodiment of the method according to the present invention can be utilized to measure the number of silica molecules (SiO₂, MW: ~60 Da) (as described in Handbook of Chemistry and Physics, 86 ed., 2005: CRC Press, p. 2544), etched by a diluted hydrofluoric acid (HF) solution. SiO₂ is a representative of small molecules, and its surface density is well known. In this example, a cover slip bottom culture dish (Mattek, Ashland, MA) was filled with de-ionized water, and the

HF solution was injected into the dish to achieve desired concentrations. The probe beam at the cover slip surface had a diameter of $\sim 5\mu\text{m}$, and the changes of the effective thickness were monitored as a function of time. Figure 9a shows a graph illustrating an exemplary change of a cover slip thickness at a particular HF concentration $\sim 0.07\%$ in volume in accordance with the present invention. For this graph of Figure 9a, the measured etching rate was $\sim 51\text{ nm/min}$. A cover slip bottom culture dish was filled with de-ionized water, and the HF solution was injected into the dish to achieve desired concentrations ($7 \times 10^{-5} \sim 0.7\%$). The change of the etching rate of the silica molecules was also measured, as varying the HF concentration, as shown in Figure 9b which illustrates a graph of an exemplary large change of an etching rate at different HF concentrations in accordance with the present invention, e.g., when the HF concentration is over 0.05% .

III. Photosynthetic protein layer imaging

The photo-synthetic proteins extracted from spinach were patterned onto a cover slip using a micro-stamp contact printing technique (as described in A. Bernard et al., "Microcontact printing of proteins," Advanced Materials, 2000, Vol. 12, pp. 1067-1070), and the pattern of the proteins was imaged with the exemplary system, arrangement and method according to the present invention, as measuring the phase in the interference between reflections from top and bottom surfaces of the cover slip. Figure 10 shows a graph 10000 of an image of a distribution of a photosynthetic protein layer generated using the arrangement and method in accordance with the present invention the surface. The thickness distribution across a cover slip was obtained by measuring phase in the interference between top and bottom surface of the cover slip. The photosynthetic protein layer was patterned by a micro-stamp contact printing technique. The result demonstrates the potential of the invention for imaging ultra thin organic layers or films.

There are several aspects of the exemplary embodiments of the system, arrangement and method according to the present invention in the implementation for chemical and biological species detection. For example, these exemplary embodiments can provide:

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- 15
- i. a label-free detection, e.g., a molecular recognition can be achieved without a specimen preparation such as fluorescence and radioactive labeling.
 - ii. the sensing area can be approximately as small as diffraction-limited size (~1 micron), and the detection can be achieved with significantly reduced amount of molecules.
 - iii. the small size of the sensing area can permit monitoring multitudes of activated probe sites in parallel on two-dimensional disposable arrays.
 - iv. the exemplary measurement system and arrangement can be completely decoupled from microarrays or microfluidic devices, and thus may be deployed to any environments, and may not use the regeneration of the sensor surface.
 - v. the multi-layer depth-resolved molecular detection can be performed.
 - vi. the measurement can be achieved at microsecond temporal resolution, and the exemplary embodiment can be applied to fast kinetic procedures such as DNA denaturization.
 - vii. the exemplary embodiment can also be used to measure the concentration and diffusion of free molecules in micro-fluidic device.

The foregoing merely illustrates the principles of the invention.

20 Various modifications and alterations to the described embodiments will be apparent to those skilled in the art in view of the teachings herein. Indeed, the arrangements, systems and methods according to the exemplary embodiments of the present invention can be used with any OCT system, OFDI system, SD-OCT system or other imaging systems, and for example with those described in International Patent

25 Application PCT/US2004/029148, filed September 8, 2004, U.S. Patent Application No. 11/266,779, filed November 2, 2005, and U.S. Patent Application No. 10/501,276, filed July 9, 2004, the disclosures of which are incorporated by reference herein in their entireties. It will thus be appreciated that those skilled in the art will be able to devise numerous systems, arrangements and methods which, although not

30 explicitly shown or described herein, embody the principles of the invention and are thus within the spirit and scope of the present invention. In addition, to the extent that the prior art knowledge has not been explicitly incorporated by reference herein

above, it is explicitly being incorporated herein in its entirety. All publications referenced herein above are incorporated herein by reference in their entireties.

WHAT IS CLAIMED IS:

1. A system comprising:
 - at least one first arrangement configured to provide a particular
5 radiation which includes at least one first electro-magnetic radiation directed to at least one sample and at least one second electro-magnetic radiation directed to a reference;
at least one second arrangement configured to detect an interference
between at least one third electro-magnetic radiation associated with the at least one
10 first electro-magnetic radiation and at least one fourth electro-magnetic radiation associated with the at least one second electro-magnetic radiation; and
at least one third arrangement configured to determine a change in a thickness of at least one portion of the at least one sample based on the interference,
wherein the particular radiation has at least one of:
15
 - i. a wavelength provided by the at least one first arrangement that varies over time, or
 - ii. a spectral width that is greater than 10nm.
2. The system according to claim 1, wherein the first and second radiations share
20 a common path.
3. The system according to claim 1, wherein the at least one sample includes a plurality of samples, and wherein the change in the thickness of the at least one portion of each of the samples is determined simultaneously.
25
4. The system according to claim 1, wherein the change in the thickness of the at least one portion of the at least one samples is determined simultaneously at different locations at least one of along or perpendicular to a beam path of the at least one first electro-magnetic radiation.
- 30 5. The system according to claim 1, wherein the change in the thickness of the at least one portion of the at least one samples is determined simultaneously along

different locations along a beam path of the at least one first electro-magnetic radiation.

6. The system according to claim 1, wherein the at least one first electro-magnetic radiation is scanned over a surface of the at least one sample at a plurality of
5 locations thereon.

7. The system according to claim 1, wherein the at least one portion of the at least one sample is coated with particular molecules that are designed to associate with or dissociate from to further molecules.

10

8. The system according to claim 7, wherein the change of the thickness is associated with an association or a dissociation of the particular molecules.

9. The system according to claim 7, wherein the particular molecules have an
15 affinity to bind to the further molecules that are different from the particular molecules.

10. The system according to claim 7, wherein the at least one portion includes a plurality of portions, wherein a first set of the particular molecules has an affinity to
20 bind to a first portion of the plurality of portions, and the a second set of the particular molecules has an affinity to bind to a second portion of the plurality of portions, and wherein the first and second sets are different from one another.

11. The system according to claim 1, wherein the at least one sample has multiple
25 layers therein.

12. The system according to claim 1, wherein the at least one sample is disposable.

13. The system according to claim 1, wherein the at least one sample is a micro-fluidic arrangement.
30

14. The system according to claim 1, wherein the change of the thickness of the at least one portion of the at least one sample is at least one of an optical thickness change, a physical thickness change or a refractive index change.
- 5 15. The system according to claim 14, wherein the thickness change is associated with a concentration of molecules at least one of on or in the at least one portion of the at least one sample.
- 10 16. The system according to claim 14, wherein the thickness change as a function of wavelength is associated with types of molecules at least one of on or in the at least one portion of the at least one sample.
- 15 17. The system according to claim 1, wherein the at least one first electromagnetic radiation has a cross-section of a beam on or in the at least one portion of the at least one sample has a size that is at least a diffraction-limited size.
18. The system according to claim 1, wherein the at least one third arrangement determines the thickness by:
- 20 i. transforming the interference into first data which is in a complex format,
 - ii. determining an absolute value associated with the first data to generate second data,
 - iii. identifying particular locations of the at least one portion as a function of the second data,
 - 25 iv. determining a phase associated with the first data to generate third data, and
 - v. associating the change of the thickness with the third data.
19. The system according to claim 1, wherein the interference is Fourier transformed to generate the first data.
- 30

20. A method comprising:
- providing a particular radiation which includes at least one first electro-magnetic radiation directed to at least one sample and at least one second electro-magnetic radiation directed to a reference;
- 5 detecting an interference between at least one third electro-magnetic radiation associated with the at least one first electro-magnetic radiation and at least one fourth electro-magnetic radiation associated with the at least one second electro-magnetic radiation; and
- determining a change in a thickness of at least one portion of the at least one
- 10 sample based on the interference,
- wherein the particular radiation has at least one of:
- i. a wavelength that varies over time, or
 - ii. a spectral width that is greater than 10nm.

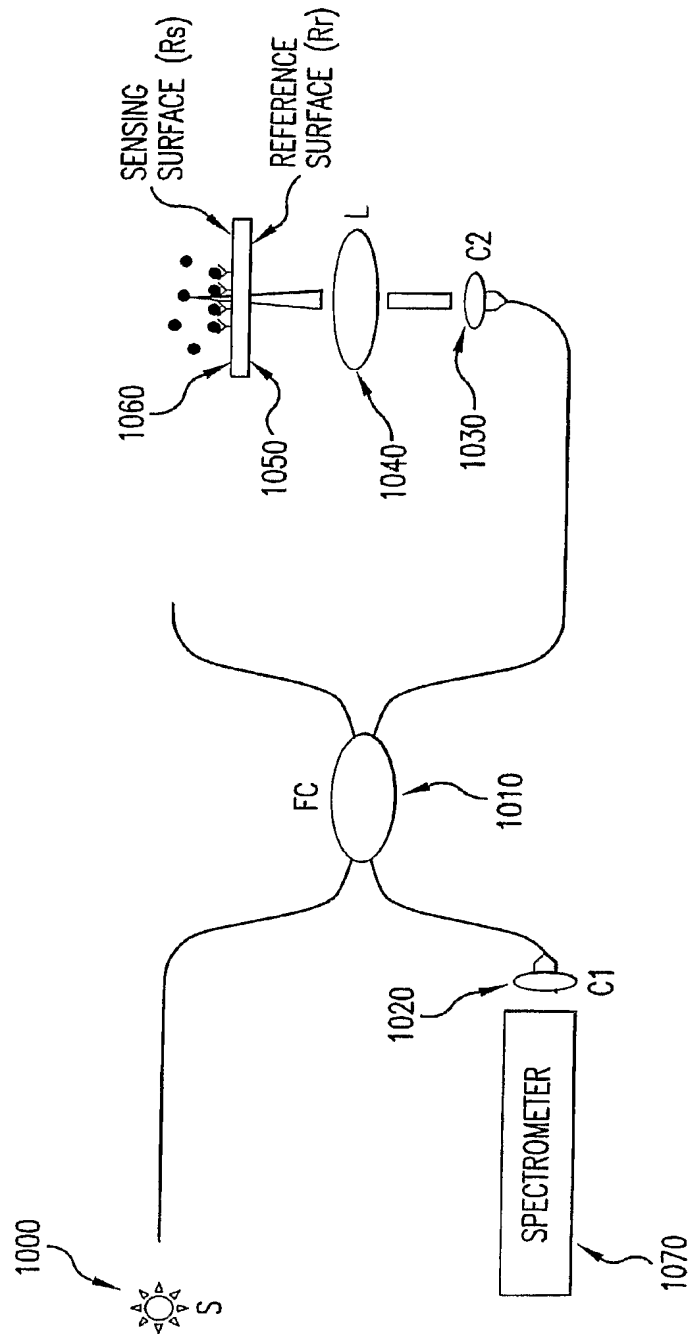
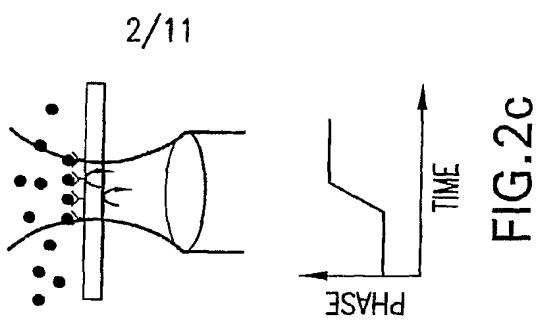
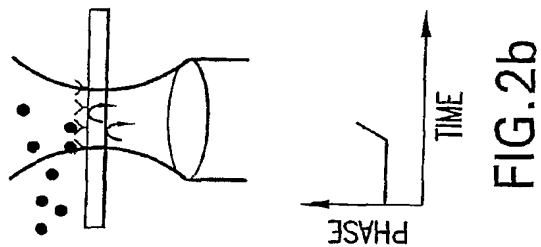
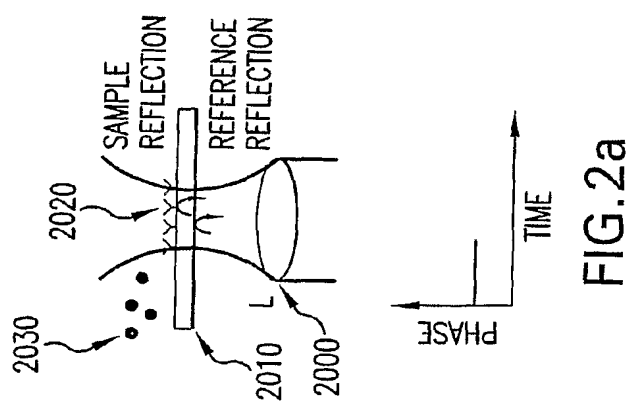


FIG. 1



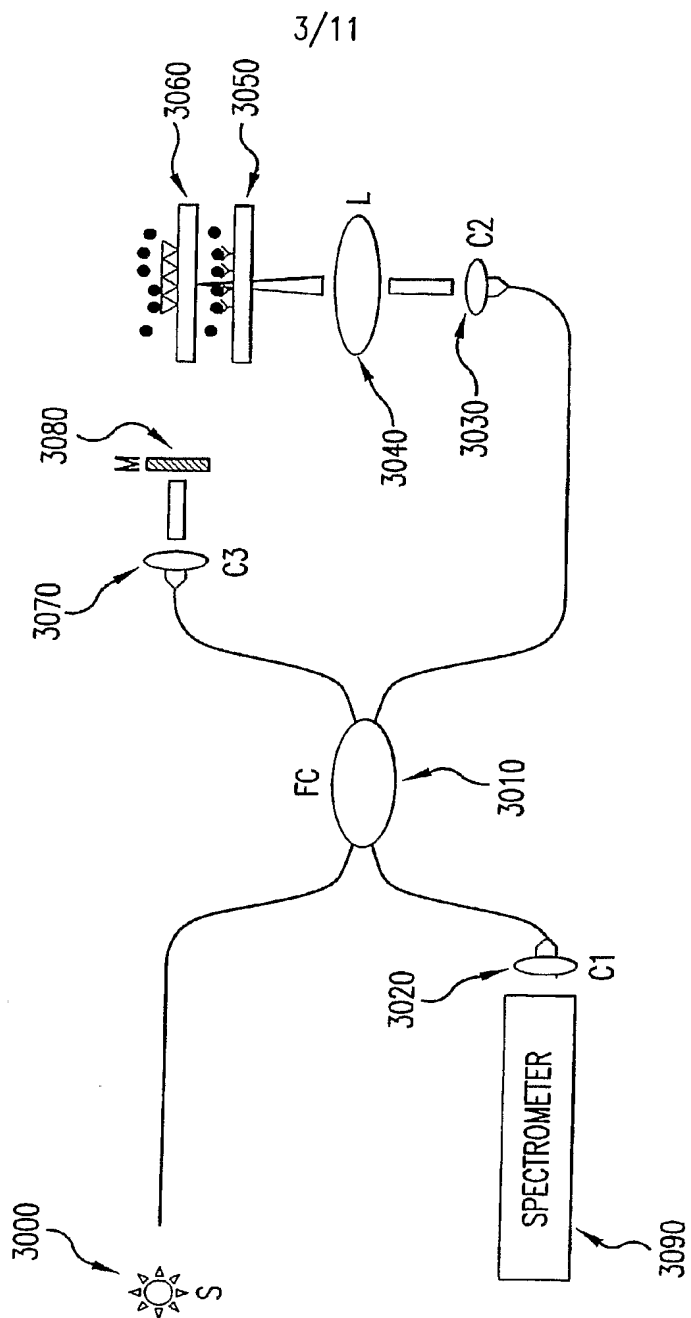


FIG.3

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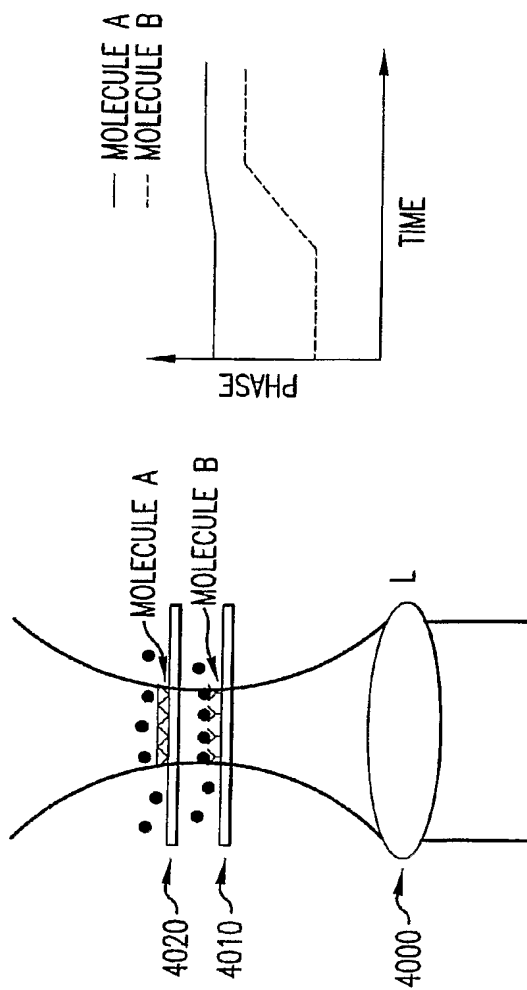


FIG.4

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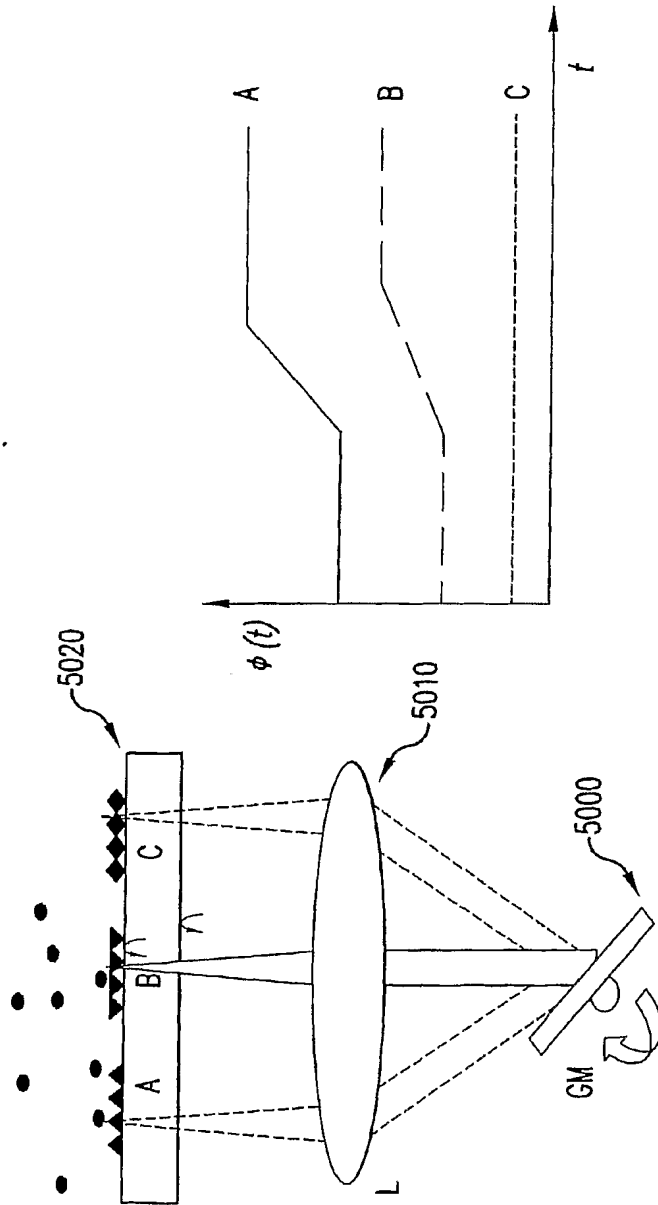
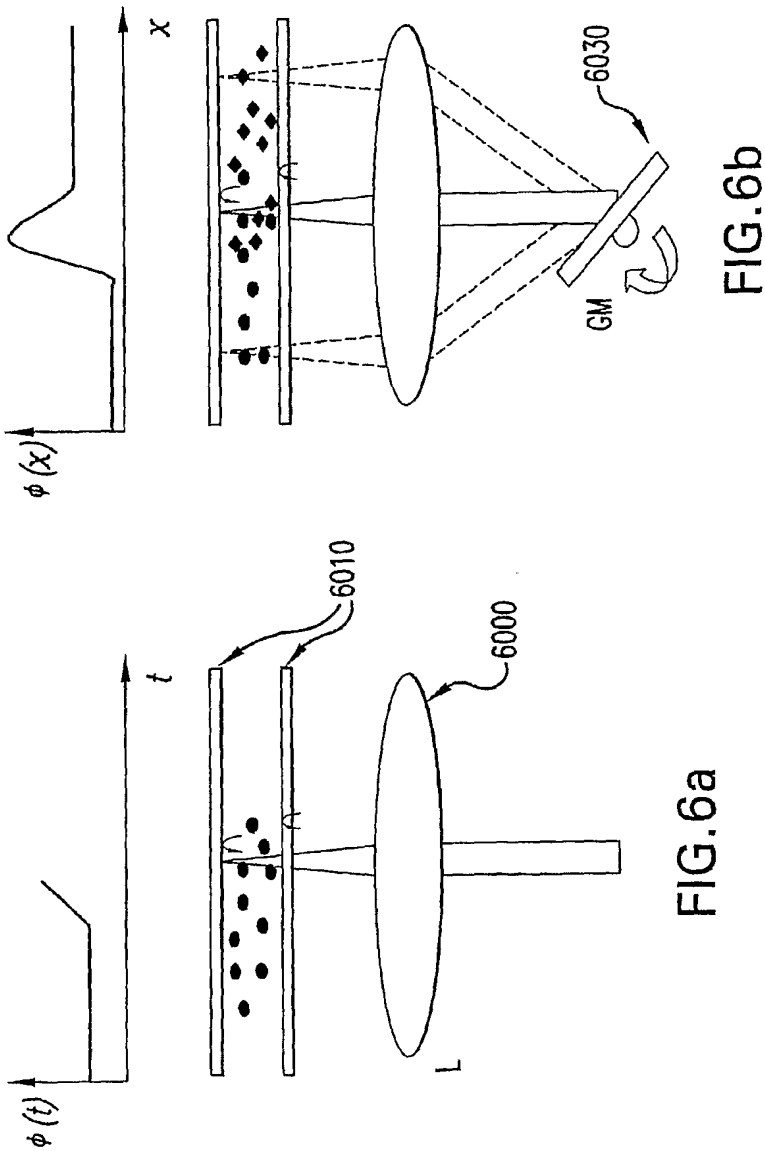


FIG.5



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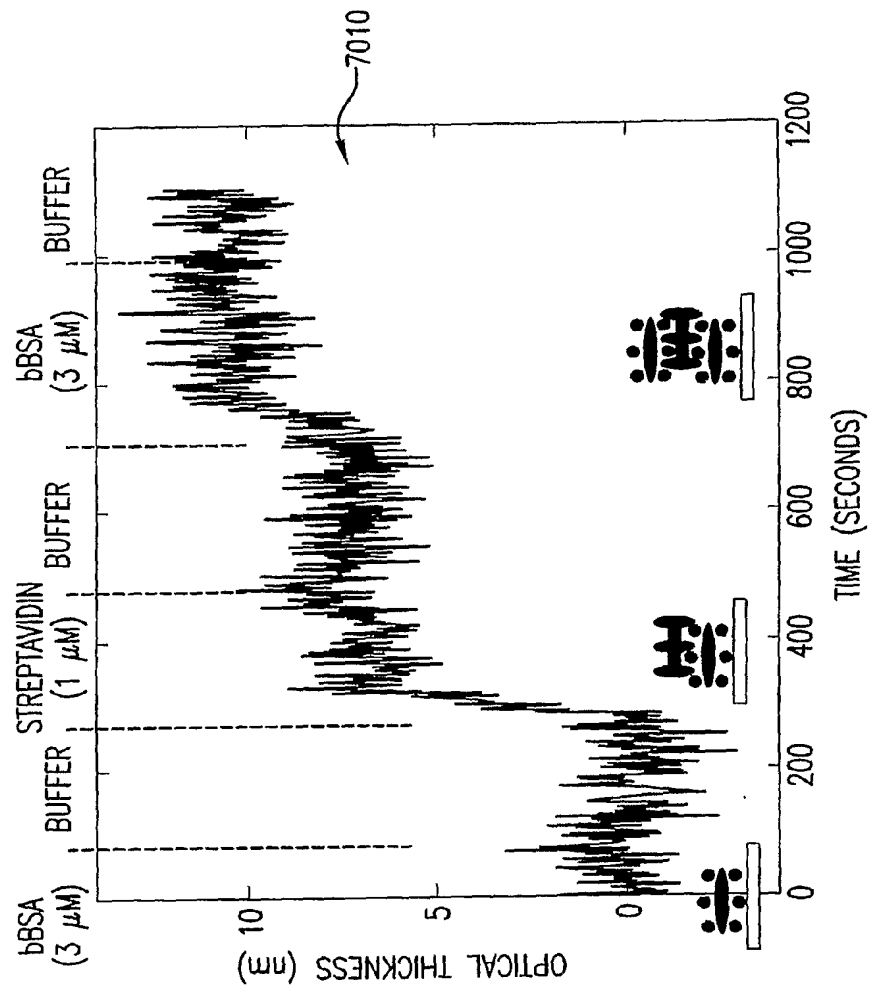


FIG. 7

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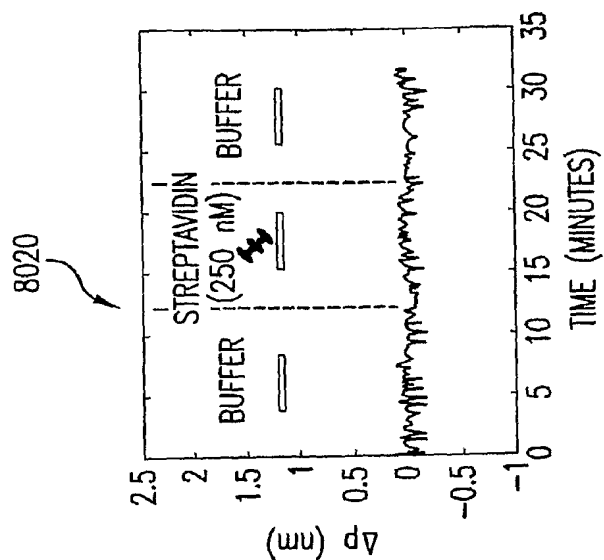


FIG. 8b

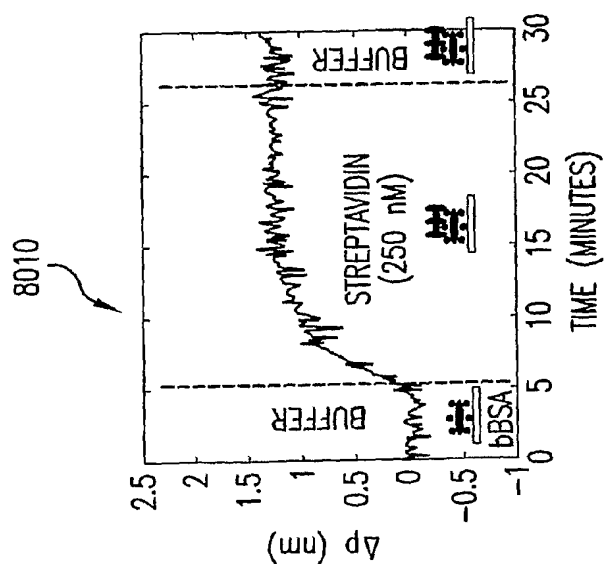


FIG. 8a

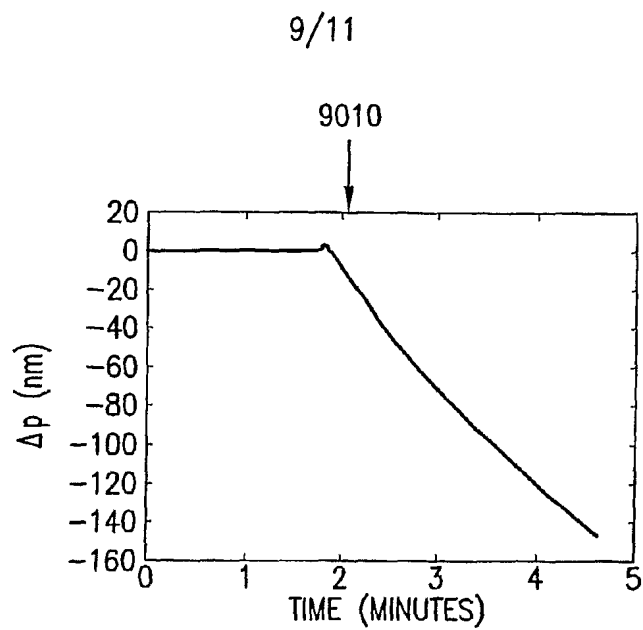


FIG. 9a

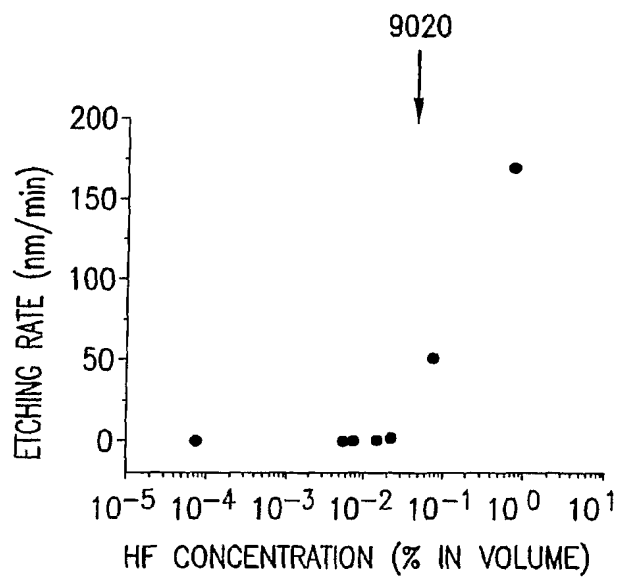


FIG. 9b

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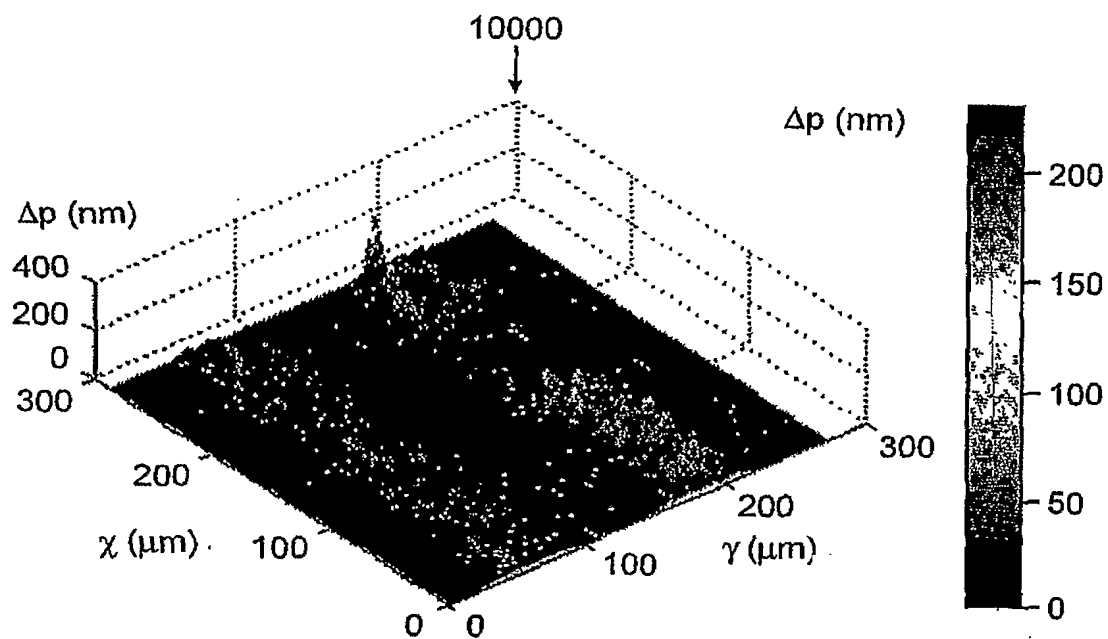


FIG.10

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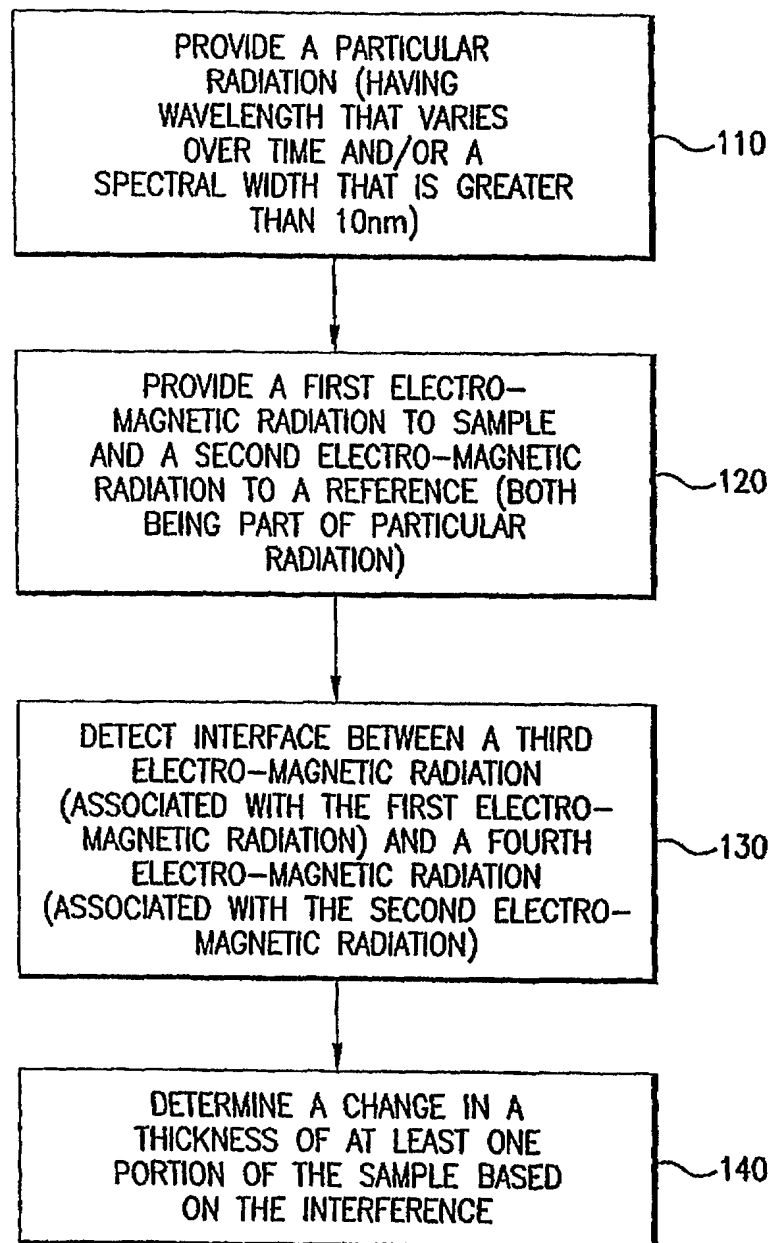


FIG.11

INTERNATIONAL SEARCH REPORT

International application No

PCT/US2006/018865

A. CLASSIFICATION OF SUBJECT MATTER INV. G01N21/47 A61B3/00 G01B9/02		
According to International Patent Classification (IPC) or to both national classification and IPC		
B. FIELDS SEARCHED Minimum documentation searched (classification system followed by classification symbols) G01N A61B G01B		
Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched		
Electronic data base consulted during the International search (name of data base and, where practical, search terms used) EPO-Internal, INSPEC, WPI Data		
C. DOCUMENTS CONSIDERED TO BE RELEVANT		
Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	B. CENSE ET AL: "Spectral domain polarization-sensitive optical coherence tomography at 850 nm" COHERENCE DOMAIN OPTICAL METHODS AND OPTICAL COHERENCE TOMOGRAPHY IN BIOMEDICINE IX 23 JAN. 2005 SAN JOSE, CA, USA, vol. 5690, no. 1, April 2005 (2005-04), pages 159-162, XP002395755 Proceedings of the SPIE - The International Society for Optical Engineering SPIE-Int. Soc. Opt. Eng USA ISSN: 0277-786X page 160 - page 161; figures 1-4 ----- -/--	1-20
<input checked="" type="checkbox"/> Further documents are listed in the continuation of Box C. <input checked="" type="checkbox"/> See patent family annex.		
* Special categories of cited documents : "A" document defining the general state of the art which is not considered to be of particular relevance "E" earlier document but published on or after the international filing date "L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified) "O" document referring to an oral disclosure, use, exhibition or other means "P" document published prior to the international filing date but later than the priority date claimed "T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention "X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone "Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art. "&" document member of the same patent family		
Date of the actual completion of the international search		Date of mailing of the international search report
23 August 2006		18/09/2006
Name and mailing address of the ISA/ European Patent Office, P.B. 5818 Patentlaan 2 NL - 2280 HV Rijswijk Tel. (+31-70) 340-2040, Tx. 31 651 epo nl, Fax: (+31-70) 340-3016		Authorized officer Consalvo, D

INTERNATIONAL SEARCH REPORT

International application No
PCT/US2006/018865

C(Continuation). DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	<p>YMETI A ET AL: "Integration of microfluidics with a four-channel integrated optical Young interferometer immunosensor"</p> <p>BIOSENSORS & BIOELECTRONICS, ELSEVIER SCIENCE PUBLISHERS, BARKING, GB, vol. 20, no. 7, 15 January 2005 (2005-01-15), pages 1417-1421, XP004761153 ISSN: 0956-5663 page 1418, column 1, line 6 - page 1420, column 2, last line ; figures 1,2</p>	1-20
A	<p>US 5 321 501 A (SWANSON ET AL) 14 June 1994 (1994-06-14) the whole document</p>	1-20
A	<p>F. LEXER ET AL: "Wavelength-tuning interferometry of intraocular distances"</p> <p>APPLIED OPTICS, vol. 36, no. 25, 1 September 1997 (1997-09-01), pages 6548-6553, XP002396007 paragraphs [004.], [005.]</p>	1-20

INTERNATIONAL SEARCH REPORT

Information on patent family members

International application No

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